Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application:

Listing of Claims:

1-50. (cancelled)

- 51. (new) A method for screening a plurality of compounds so as to identify at least one compound exhibiting cognitive enhancing activity, comprising:
 - a) determining in vitro efficacy and EC50 values for each compound at an $\alpha_1\beta_2\gamma_2$ or an $\alpha_5\beta_3\gamma_2$ GABAA subtype receptor;
 - b) determining an in vitro efficacy value for each compound at a GABAA receptor comprising an α_2 or α_3 subunit; and
 - c) identifying as exhibiting cognitive enhancing activity a compound having: an EC_{50} value determined in a) of less than about 200nM, an efficacy value determined in a) of less than about -5%, and an efficacy value determined in b) of greater than about 5%.
- 52. (new) The method of Claim 51 wherein the EC_{50} measured in step a) is less than 150 nM.

- 53. (new) The method of Claim 52 wherein the *in vitro* efficacy measured at said $\alpha_1\beta_2\gamma_2$ GABA_A subtype receptor or said $\alpha_5\beta_3\gamma_2$ GABA_A subtype receptor is less than -10%.
- 54. (new) The method of Claim 53 wherein the *in vitro* efficacy measured at said GABA_A receptor comprised of said α_2 subunit or said α_3 subunit is greater than 10%.
- 55. (new) The method of Claim 51 wherein the *in vitro* efficacy measured at said $\alpha_1\beta_2\gamma_2$ GABA_A subtype receptor or said $\alpha_5\beta_3\gamma_2$ GABA_A subtype receptor is less than -10%.
- 56. (new) The method of Claim 55 wherein the *in vitro* efficacy measured at said GABA_A receptor comprised of said α_2 or said α_3 subunit is greater than 10%.
- 57. (new) The method of Claim 51 wherein the GABA_A receptor comprised of said α_2 subunit is an $\alpha_2\beta_3\gamma_2$ GABA_A receptor or the GABA_A receptor comprised of said α_3 subunit is an $\alpha_3\beta_3\gamma_2$ GABA_A receptor.
- 58. (new) A method for screening compounds for cognitive enhancing activity, comprising:
 - a) selecting compounds having a binding affinity less than 100 nM at any $GABA_A$ receptor;
 - b) determining in vitro efficacy and EC50 values for each selected compound at an $\alpha_1\beta_2\gamma_2$ or $\alpha_5\beta_3\gamma_2$ GABA, subtype receptor;

- c) determining in vitro efficacy and EC50 values for each selected compound at a GABAA receptor comprised of an α_2 or α_3 subunit; and
- d) identifying as having cognitive enhancing activity any compound having an EC_{50} value determined in b) of less than 200nM and an efficacy value measured in b) of less than -5%, and an efficacy value measured in c) of greater than 5%.
- 59. (new) A method for screening a plurality of compounds for cognitive enhancing activity, comprising:
 - a) determining in vitro efficacy and EC50 values for each compound at $\alpha_1\beta_2\gamma_2$ or $\alpha_5\beta_3\gamma_2$ GABAA receptors;
 - b) determining in vitro efficacy for each compound at a GABAA receptor comprised of an α_2 or α_3 subunit;
 - c) determining the *in vivo* effect of each compound in an animal model for measuring cognitive enhancement;
 - d) determining the *in vivo* effects of each compound in an animal model for proconvulsant activity by measuring a seizure threshold in the presence of a seizure inducing compound or in an animal model that predicts anxiogenic effects; and
 - e) identifying a cognitive enhancing compound as a compound having cognitive enhancing properties when the EC_{50} measured in step a) is less than 200nM and the efficacy measured in step a) is less than -5% and the efficacy measured in step b) is greater than 5% and said compound produces a statistically significant (p <0.05) positive effect in the animal model indicative of cognitive enhancement and said compound does not produce an effect in the animal model predictive of

proconvulsant activity of more than a 25% decrease in the seizure threshold in the presence of the seizure inducing drug, or does not produce a change that is statistically significant in said model, or the compound does not produce a statistically significant effect in the animal model that predicts anxiogenic effects.

- 60. (new) A method for screening compounds for cognitive enhancing properties, comprising:
 - a) selecting compounds having binding affinities of less than 100 nM at any GABA_A receptor;
 - b) measuring the in vitro efficacy of each compound at an $\alpha_1\beta_2\gamma_2$ or $\alpha_5\beta_3\gamma_2$ GABAA receptor;
 - c) measuring the *in vitro* efficacy of each compound at a GABA_A receptor comprised of an α_2 or α_3 subunit;
 - d) measuring the *in vivo* effect of each compound in an animal model predictive of cognitive enhancement;
 - e) measuring the *in vivo* side effects of each compound in an animal model that predicts proconvulsant activity by measuring a seizure threshold in the presence of a seizure inducing compound or measuring the *in vivo* side effects of each compound in an animal model that predicts anxiogenic effects; and
 - f) identifying as a cognitive enhancing compound a particular compound for which the EC_{50} measured in step b) is less than 200nM and the efficacy measured in
 - step b) is less than -5% and the efficacy measured in step c) is greater than 5% and said particular compound produces a statistically significant (p
 - < 0.05) positive effect in the animal model indicative

of cognitive enhancement and said particular compound does not produce an effect in the animal model predictive of proconvulsant activity of more than a 25% decrease in the seizure threshold in the presence of the seizure inducing drug, or does not produce a change that is statistically significant in said model, or said particular compound does not produce a statistically significant effect in the animal model that predicts anxiogenic effects.

- 61. (new) A method for screening a plurality of compounds so as to identify at least one compound exhibiting cognitive enhancing activity, comprising:
 - a) determining in vitro efficacy and EC $_{50}$ values for each compound at an $\alpha_1\beta_2\gamma_2$ and an $\alpha_5\beta_3\gamma_2$ GABA, subtype receptor;
 - b) determining an in vitro efficacy value for each compound at a GABA receptor comprising an α_2 or α_3 subunit; and
 - c) identifying as exhibiting cognitive enhancing activity a compound having: EC_{50} values determined in a) of less than about 200nM at each subtype receptor, efficacy values determined in a) of less than about -5% at each subtype receptor, and an efficacy value determined in b) of greater than about 5%.
- 62. (new) The method of Claim 61 wherein the EC_{50} values measured in step a) are less than 150 nM at each subtype receptor.

- 63. (new) The method of Claim 62 wherein the $in\ vitro$ efficacy values measured in step a) are less than -10% at each subtype receptor.
- 64. (new) The method of Claim 63 wherein the *in vitro* efficacy measured at said GABA_A receptor comprised of said α_2 subunit or said α_3 subunit is greater than 10%.
- 65. (new) The method of Claim 61 wherein the *in vitro* efficacy values measured in step a) are less than -10% at each subtype receptor.
- 66. (new) The method of Claim 65 wherein the *in vitro* efficacy measured at said GABA_A receptor comprised of said α_2 or said α_3 subunit is greater than 10%.
- 67. (new) The method of Claim 61 wherein the GABA_A receptor comprised of said α_2 subunit is an $\alpha_2\beta_3\gamma_2$ GABA_A receptor or the GABA_A receptor comprised of said α_3 subunit is an $\alpha_3\beta_3\gamma_2$ GABA_A receptor.
- 68. (new) A method for screening a plurality of compounds so as to identify at least one compound exhibiting cognitive enhancing activity, comprising:
 - a) determining in vitro efficacy and EC50 values for each compound at an $\alpha_1\beta_2\gamma_2$ and an $\alpha_5\beta_3\gamma_2$ GABAA subtype receptor;
 - b) determining an in vitro efficacy value for each compound at a GABAA receptor comprising an α_2 or α_3 subunit; and

- c) identifying as exhibiting cognitive enhancing activity a compound having: EC_{50} values determined in a) of less than about 200nM at each subtype receptor, an efficacy value determined in a) of less than about -10% at the $\alpha_5\beta_3\gamma_2$ GABAA subtype receptor, an efficacy value determined in a) of greater than about 10% at the $\alpha_1\beta_2\gamma_2$ GABAA subtype receptor, and an efficacy value determined in b) of greater than about 5%.
- 69. (new) The method of Claim 68 wherein the EC_{50} values measured in step a) are less than 150 nM at each subtype receptor.
- 70. (new) The method of Claim 68 wherein the *in vitro* efficacy measured at said GABA_A receptor comprised of said α_2 subunit or said α_3 subunit is greater than 10%.
- 71. (new) The method of Claim 68 wherein the GABA_A receptor comprised of said α_2 subunit is an $\alpha_2\beta_3\gamma_2$ GABA_A receptor or the GABA_A receptor comprised of said α_3 subunit is an $\alpha_3\beta_3\gamma_2$ GABA_A receptor.